

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

IN RE APPLICATION OF

SCHURMAN ET AL.

APPLICATION NO:

FILED:

FOR: COMBINATIONS OF IMMUNOSUPPRESSIVE AGENTS FOR THE  
TREATMENT OR PREVENTION OF GRAFT REJECTIONS

Assistant Commissioner for Patents  
Washington, D.C. 20231

PRELIMINARY AMENDMENT

Sir:

Prior to the examination of the above-referenced patent application, please amend the application as follows:

**In the Claims:**

**Please cancel claim 4.**

**Please amend claims 5-6 as follows:**

5. (Amended) A pharmaceutical composition according to claim 1 wherein the pharmaceutically acceptable salt of mycophenolic acid is MPA sodium salt formulated as an enteric-coated solid oral dosage form.
6. (Amended) A method for the treatment or prevention of xenograft rejection comprising administering a pharmaceutical composition according to claim 1.

**Please add claims 10 and 11 as follows:**

10. A pharmaceutical composition according to claim 2 wherein the pharmaceutically acceptable salt of mycophenolic acid is MPA sodium salt formulated as an enteric-coated solid oral dosage form.
11. A method for the treatment or prevention of xenograft rejection comprising administering a pharmaceutical composition according to claim 2.

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**REMARKS**

Claims 5-6 have been amended, claims 10-11 have been added and claim 4 has been canceled. Favorable consideration of this application is respectfully requested.

Attached hereto is a marked-up version of the changes made to the specification and claims by the current amendment. The attached pages are captioned "VERSIONS WITH MARKINGS TO SHOW CHANGES MADE."

Respectfully submitted,

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Date: *November 7, 2001*

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VERSION WITH MARKINGS TO SHOW CHANGES MADE

In the Claims:

Claims 5-6 have been amended as follows:

5. A pharmaceutical composition according to claim 1 ~~or~~ 2 wherein the pharmaceutically acceptable salt of mycophenolic acid is MPA sodium salt formulated as an enteric-coated solid oral dosage form.
6. A method for the treatment or prevention of xenograft rejection comprising administering a pharmaceutical composition according to claim 1 ~~or~~ 2.